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Febrivac 3-Plus injektioneste, suspensio

Authorised

- Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
- Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
- Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
- Clostridium botulinum, type C, toxoid
- Mink enteritis virus, strain E-MINK F1, Inactivated

Product identification

Medicine name:
Febrivac 3-Plus injektioneste, suspensio

Active substance:
Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated

Clostridium botulinum, type C, toxoid

Mink enteritis virus, strain E-MINK F1, Inactivated

Target species:
Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
8.00 log10cell count / 1.00 millilitre(s)

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
8.00 log10cell count / 1.00 millilitre(s)

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
8.00 log10cell count / 1.00 millilitre(s)

Clostridium botulinum, type C, toxoid
0.50 unit(s) / 1.00 millilitre(s)

Mink enteritis virus, strain E-MINK F1, Inactivated
4.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Package description:

500 ml vial
250 ml vial
50 ml vial

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
CZ Vaccines S.A.U.

Marketing authorisation date:
10/12/1996

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
Finnish Medicines Agency

Authorisation number:
13860

Date of authorisation status change:
10/12/1996

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.